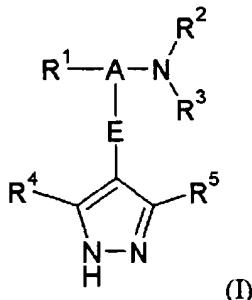


ABSTRACTPHARMACEUTICAL COMPOUNDS

The invention provides compounds of the formula (I) having protein kinase B inhibiting activity:



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wherein A is a saturated hydrocarbon linker group containing from 1 to 7 carbon atoms, the linker group having a maximum chain length of 5 atoms extending between R¹ and NR²R³ and a maximum chain length of 4 atoms extending between E and NR²R³, wherein one of the carbon atoms in the linker group may optionally be replaced by an oxygen or nitrogen atom; and wherein the carbon atoms of the linker group A may optionally bear one or more substituents selected from oxo, fluorine and hydroxy, provided that the hydroxy group when present is not located at a carbon atom α with respect to the NR²R³ group and provided that the oxo group when present is located at a carbon atom α with respect to the NR²R³ group;

10 E is a monocyclic or bicyclic carbocyclic or heterocyclic group;

15 R¹ is an aryl or heteroaryl group; and

R², R³, R⁴ and R⁵ are as defined in the claims.

Also provided are pharmaceutical compositions containing the compounds, methods for preparing the compounds and their use as anticancer agents.